Attorney's Docket No.: 06275-0472US1 / 101017-1P US/R&I

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Amendments to the Claims:

This listing of claims replaces all prior versions and listings of claims in the application:

Listing of Claims:

 (Currently Amended) A compound of formula (I) or a pharmaceutically acceptable salt thereof:

in which:

(I)

X is C₁₋₆alkyl or OR⁶;

Y is selected from hydrogen, halogen, CN, nitro, SO₂R³, OR⁴, SR⁴, SOR²; SO₂NR⁴R⁵, CONR⁴R⁵, NR⁴SO₂R³, NR⁶CO₂R⁶, NR⁶COR³, C₂·C₆ alkenyl, C₂·C₆ alkenyl, C₂·C₆ alkenyl, C₂·C₇ eyeloalkyl or C₁₋₆alkyl, the latter four groups being optionally substituted by one or more substitutents independently selected from halogen, OR⁶ and NR⁶R², S(O)₆R⁶; n is 0, 1 or 2;

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Z is phenyl optionally substituted by one or more substituents independently selected from hydrogen, halogen, CN, OH, SH, nitro, COR⁹, CO₂R⁶, SO₂R⁹, OR⁹, SR⁹, SOR⁹, SO₂NR¹⁰R¹¹, CONR¹⁰R¹¹, NHSO₂R⁹, NR⁹SO₂R⁹, NR⁶CO₂R⁶, NHCOR⁹, NR⁹COR⁹, NR⁶CONR⁴R⁵, NR⁶SO₂NR⁴R⁵, arvl.

$$\begin{split} &C_2\text{--}C_6 \text{ alkenyl, } C_2\text{--}C_6 \text{ alkynyl, } C_3\text{--}C_7 \text{ cycloalkyl or } C_{1\text{--}6}\text{alkyl, the latter four groups being} \\ &\text{optionally substituted by one or more substituents independently selected from halogen, } C_3\text{--}C_7 \text{ cycloalkyl, } OR^6, NR^6R^7, S(O)_8R^6, CONR^6R^7, NR^6COR^7, SO_2NR^6R^7 \text{ and } NR^6SO_2R^7. \end{split}$$

 R^1 and R^2 independently represent a hydrogen atom, halogen, C_2 - C_6 alkenyl, C_2 - C_6 alkenyl, C_3 - C_7 cycloalkyl or a C_{1-6} alkyl group, the latter four groups being optionally substituted by one or more substituents independently selected from halogen,

C3-C7 cycloalkyl, NR6R7, OR6, S(O)nR6;

 R^3 represents C_3 - C_7 cycloalkyl or C_{1-6} alkyl which may be optionally substituted by one or more substituents independently selected from halogen, C_3 - C_7 cycloalkyl, OR^6 and NR^6R^7 , $S(O)_nR^6$, $CONR^6R^7$, NR^6COR^7 , $SO_2NR^6R^7$ and $NR^6SO_2R^7$;

R⁴ and R⁵ independently represent hydrogen, C₃-C₇ cycloalkyl or C₁₋₆alkyl, the latter two groups being optionally substituted by one or more substituents independently selected from halogen, C₃-C₇ cycloalkyl, OR⁶ and NR⁶R⁷, S(O)_nR⁶, CONR⁶R⁷, NR⁶COR⁷,SO₂NR⁶R⁷ and NR⁶SO₂R⁷;

R⁶ and R⁷ independently represents a hydrogen atom or C₁-C₆ alkyl;

 $R^8 \text{ is hydrogen, } C_{1\text{--}4} \text{ alkyl, -COC}_1\text{--}C_4 \text{ alkyl, } CO_2C_1\text{--}C_4 \text{alkyl or } CONR^6C_1\text{--}C_4 \text{alkyl; } \\$

 R^9 represents aryl, C_3 - C_7 cycloalkyl or $C_{1.6}$ alkyl, the latter two groups may be optionally substituted by one or more substituents independently selected from halogen, C_3 - C_7 cycloalkyl, aryl, OR^6 and NR^6R^7 , $S(O)_nR^6$, $CONR^6R^7$, NR^6COR^7 , $SO_2NR^6R^7$ and $NR^6SO_2R^7$;

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R¹⁰ and R¹¹ independently represent aryl, hydrogen, C₃-C₇ cycloalkyl or C₁₋₆alkyl, the latter two groups being optionally substituted by one or more substituents independently selected from halogen, C₃-C₇ cycloalkyl, aryl, OR⁶ and NR⁶R⁷, S(O)_nR⁶, CONR⁶R⁷, NR⁶COR⁷, SO₂NR⁶R⁷ and NR⁶SO₂R⁷.

- 2. (Previously Presented) A compound according to claim 1 in which R^1 and R^2 independently represent a hydrogen atom, C_2 - C_6 alkenyl, C_2 - C_6 alkenyl, C_3 - C_7 cycloalkyl or a C_1 -ealkyl group, the latter four groups being optionally substituted by one or more substituents independently selected from halogen, C_3 - C_7 cycloalkyl, NR^6R^7 , OR^6 , $S(O)_nR^6$.
- (Previously presented) A compound according to claim 1 in which X is C₁₋₄alkyl or C₁₋₄alkoxy.
- 4. (Cancelled)
- (Cancelled)
- (Previously Presented) A compound according to claim 1 in which Z is substituted by one or more substituents independently selected from halogen, C_{1,3}alkyl, evano and SO₂R⁹.
- (Previously presented) A compound according to claim 1 in which R¹ and R² are both hydrogen or one is hydrogen and the other is C₁₋₃ alkyl.
- 8. (Currently Amended) A compound according to claim 1 selected from: [(5-Methylbiphenyl-2-yl)oxy]acetic acid, {[5-Ethyl-4'-(methylsulfonyl)biphenyl-2-yl]oxy}acetic acid,

{[4'-(Ethylsulfonyl)-5-methoxybiphenyl-2-ylloxy}acetic acid,

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[[4 Chloro 4' (ethylsulfonyl) 2',5-dimethyl[1,1'-biphenyl] 2-yl]oxy]-acetic acid, [[4'-(Ethylsulfonyl)-2',5-dimethyl[1,1'-biphenyl]-2-yl]oxy]-acetic acid, 2-[[3'-Cyano-5-methyl[1,1'-biphenyl]-2-yl]oxy]-(2S)-propanoic acid, 2-[[2'-Fluoro-5'-cyano-5-methyl[1,1'-biphenyl]-2-yl]oxy]-(2S)-propanoic acid, and pharmaceutically acceptable salts thereof.

Claims 9-11 (Cancelled)

- 12. (Previously presented) A method for the therapeutic treatment of asthma or rhinitis in a patient suffering from asthma or rhinitis, which comprises administering to the a patient suffering from asthma or rhinitis a therapeutically effective amount of a compound of formula (I), or a pharmaceutically acceptable salt as defined in claim 1.
- (Previously presented) A compound according to claim 2 in which X is C₁₋₄alkyl or C₁₋₄alkoxy.
- (Cancelled)
- (Cancelled)
- (Previously presented) A compound according to claim 2 in which Z substituted by one
 or more substituents independently selected from halogen, C₁₋₃alkyl, cyano and SO₂R⁹.
- 17. (Previously presented) A compound according to claim 2 in which R^1 and R^2 are both hydrogen or one is hydrogen and the other is $C_{1:3}$ alkyl.

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18. (Previously presented) A pharmaceutical composition comprising a compound of formula (I) as claimed in claim 1, or a pharmaceutically acceptable salt thereof, and a pharmaceutically acceptable adjuvant, diluent, or carrier.

19. (Previously presented) A method of producing a CRTh2 receptor inhibitory effect in a patient, which comprises administering to the patient an effective amount of a compound of formula (I) as claimed in claim 1 or a pharmaceutically acceptable salt thereof.